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AMENDMENTS TO THE CLAIMS

Please amend claims 1, 4, 9-10, 13-21, and 23-24 as indicated below.

Please cancel claim 11 without prejudice.

The listing of claims will replace all prior versions, and listings, of claims in the application.

Listing of Claims:

1. (Currently Amended) An isolated compound having of the structure (I):

$$(R_4)_x$$
 R_3
 E_4
 R_1
 R_2
 R_1

wherein:

R₁ is a substituted alkyl;

R₂ is methyl;

R₃ is hydroxy;

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Each R₄ is independently alkyl, substituted alkyl, alkenyl, substituted alkynyl, substituted alkynyl, aryl, substituted aryl, cycloalkyl, or substituted cycloalkyl;

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 E_1 , E_3 , and E_4 are-O;

 E_2 is $-NR_5$, wherein R_5 is -H or C_1 - C_6 alkyl; and

X is 0 to 8.

- 2. (Previously Presented) The compound of Claim 1, wherein E2 is -NH.
- 3. (Canceled).
- 4. (Currently Amended) The compound of Claim 1, wherein R₁ is a substituted alkyl substituted with one or more substitutions substituents selected from the group consisting of halogen, cyano, oxyacyl, amino, amido, -C(O)H, acyl, carboxyl, and sulfonamide.
- 5. (Original) The compound of Claim 4, wherein the substituted alkyl is a halogenated alkyl.
- 6. (Original) The compound of Claim 5, wherein the halogenated alkyl is a chlorinated alkyl.
- 7. (Previously Presented) A pharmaceutical composition comprising at least one compound of Claim 1 in a pharmaceutically acceptable carrier thereof.
- 8. (Canceled).
- 9. (Currently Amended) The pharmaceutical composition of Claim 7, further comprising at least one additional anti-neoplastic agent in combination with the at least one compound of claim 1.

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10. (Currently Amended) A method of treating a <u>human</u> refractile <u>mammalian cancer</u> cell <u>proliferative disorder</u>, comprising <u>administering to a subject in need thereof contacting human refractile cancer cells with a therapeutically effective amount of a compound <u>having of</u> the structure (I):</u>

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$$(R_4)_x$$
 R_3
 E_4
 R_1
 R_2
 R_1
 R_2

wherein:

R₁ to R₃ are each independently -H, alkyl, substituted alkyl, alkenyl, substituted alkenyl, alkynyl, substituted alkynyl, aryl, substituted aryl, heteroaryl, substituted heteroaryl, heterocyclic, substituted heterocyclic, cycloalkyl, substituted cycloalkyl, alkoxy, substituted alkoxy, thioalkyl, substituted thioalkyl, hydroxy, halogen, amino, amido, carboxyl, -C(O)H, acyl, oxyacyl, carbamate, sulfonamide, or sulfuryl;

Each R₄ is independently alkyl, substituted alkyl, alkenyl, substituted alkynyl, substituted alkynyl, aryl, substituted aryl, cycloalkyl, <u>or</u> substituted cycloalkyl;

 E_1 to E_4 are each independently -O,- NR_5 , or -S, wherein R_5 is -H or C_1 - C_6 alkyl; and

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X is 0 to 8.

- 11. (Canceled).
- 12. (Currently Amended) The method of Claim 10, wherein the compound has is of the structure of formula (V):

13. (Currently Amended) The method of Claim 10, wherein the <a href="https://human.com

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14. (Currently Amended) The method of Claim 13, wherein the human refractile cancer cell proliferative disorder is a sarcoma eell of a soft tissue or a sarcoma eell of a bone.

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15. (Currently Amended) The method of Claim 13, wherein the human refractile cancer cell proliferative disorder is a leukemia cancer cell.

- 16. (Currently Amended) The method of Claim 13, wherein the human refractile cancer cell proliferative disorder is a myeloma cancer cell.
- 17. (Currently Amended) The method of Claim 13, wherein the human refractile cancer cell proliferative disorder is an ovarian cancer cell.
- 18. (Currently Amended) The method of Claim 13, wherein the human refractile cancer cell proliferative disorder is a prostate cancer cell.
- 19. (Currently Amended) The method of Claim 13, wherein the human refractile cancer cell proliferative disorder is a non-Hodgkin's disease cancer cell.
- 20. (Currently Amended) The method of Claim 13, wherein the human refractile eaneer cell proliferative disorder is a pancreatic adenocarcinoma eancer cell.
- 21. (Currently Amended) The method of Claim 10, wherein: R₁ is a substituted alkyl; R₂ is methyl; R₃ is hydroxy; each R₄ is independently alkyl, substituted alkyl, alkenyl, substituted alkenyl, alkynyl, substituted alkynyl, aryl, substituted aryl, cycloalkyl, or substituted cycloalkyl; E₁, E₄ and E₃ are -O; E₂ is -NR₅, wherein R₅ is -H or C₁-C₆ alkyl; and X is 0 to 8.
- 22. (Previously Presented) The method of Claim 21, wherein R₁ is a halogenated alkyl.
- 23. (Currently Amended) The method of Claim 10, further comprising administering at least one additional anti-neoplastic agent in combination with at least one compound of the structure (I).

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24. (Currently Amended) A pharmaceutical composition comprising a pharmaceutically acceptable carrier and an effective amount of a compound having of the structure:

wherein the pharmaceutical composition is in a solid form.

25. (Previously Presented) The compound of Claim 1, wherein the structure is: